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# Three-dimensional broadband tunable terahertz metamaterials

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We present optically tunable magnetic three-dimensional (3D) metamaterials at terahertz (THz) frequencies which exhibit a tuning range of  $\sim 30\%$  of the resonance frequency. This is accomplished by fabricating 3D array structures consisting of double-split-ring resonators (DSRRs) on silicon on sapphire, fabricated using multilayer electroplating. Photoexcitation of free carriers in the silicon within the capacitive region of the DSRR results in a redshift of the resonant frequency from 1.74 to 1.16 THz. The observed frequency shift leads to a transition from a magnetic-to-bianisotropic response as verified through electromagnetic simulations and parameter retrieval. Our approach extends dynamic metamaterial tuning to magnetic control, and may find applications in switching and modulation, polarization control, or tunable perfect absorbers.

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The advent of artificial electromagnetic materials has provided unique routes to engineer photonic media, leading to the realization of intriguing phenomena. This includes the successful demonstration of invisibility cloaking<sup>1–3</sup> and negative refraction.<sup>4,5</sup> Scale invariance of the underlying equations enables translation of phenomena realized in one region of the electromagnetic spectrum to others. Metamaterials and, in particular, split-ring resonators (SRRs), can couple to the electric, magnetic, or both fields of incident electromagnetic radiation. This leads to a Lorentzian-like effective response in the permittivity, permeability, and bianisotropy, described by the general constitutive relations as

$$\begin{pmatrix} \mathbf{p} \\ \mathbf{m} \end{pmatrix} = \begin{pmatrix} \alpha_{EE} & \alpha_{EH} \\ \alpha_{HE} & \alpha_{HH} \end{pmatrix} \begin{pmatrix} \mathbf{E}_{\text{in}} \\ \mathbf{H}_{\text{in}} \end{pmatrix}, \quad (1)$$

where  $\alpha_{EE}$  ( $\alpha_{HH}$ ) is the tensorial purely electric (magnetic) polarizability, quantifying the electric (magnetic) dipole induced purely by a incident electric (magnetic) field and the off-diagonal  $\alpha_{EH}$  ( $\alpha_{HE}$ ) is the tensorial magnetoelectric coupled polarizability, quantifying the electric (magnetic) dipole induced by an incident magnetic (electric) field. The Onsager relations require  $\alpha_{EH} = -\alpha_{HE}^T$ .<sup>6</sup> The bianisotropic parameter  $\xi$  relates to the magnetoelectric polarizability by  $\xi = i c_0 \alpha_{EH}$ , where  $c_0$  is the velocity of light in free space.

By tailoring the geometry or configuration of subwavelength metallic unit cells, the effective parameters can be specified and controlled, leading to myriad routes to control the amplitude, frequency, and phase of the incident electromagnetic radiation. For example, by independently tuning the electric and magnetic resonance of multilayer composites, it is possible to match the impedance to free space forming a perfect absorber with minimized reflectance.<sup>7–9</sup> This ability to manipulate incident radiation is particularly important at terahertz frequencies given the relative paucity of devices. Recent progress on tunable metamaterial through voltage,<sup>10,11</sup> optical,<sup>12–14</sup> thermal,<sup>15–17</sup> and mechanical<sup>18,19</sup> tuning, has shown its potential to fill the “THz gap.”

The majority of previous studies are based on planar metamaterials, and therefore result in tuning the electric response. It is desirable to explore new configurations to

access magnetic control of metamaterials which will, in turn, provide additional flexibility in tailoring the electromagnetic response for potential applications such as tunable absorbers or polarizers. Recently, three-dimensional metamaterials at terahertz frequencies have been realized with the successful demonstration of negative refraction<sup>20</sup> and negative permeability,<sup>21</sup> with one example of a tunable three-dimensional (3D) metamaterial.<sup>22</sup> We demonstrate optically tunable magnetic 3D metamaterials exhibiting a tuning range of  $\sim 30\%$  of the resonance frequency. This is accomplished by fabricating 3D array structures consisting of double-split-ring resonators (DSRRs) on silicon on sapphire, fabricated using multilayer electroplating. Photoexcitation of free carriers in the silicon within the capacitive region of the DSRR results in a redshift of the resonant frequency from 1.74 to 1.16 THz. Further, the observed frequency shift results in a transition from a magnetic-to-bianisotropic response as verified through electromagnetic simulations and parameter retrieval.

The tuning principle and unit cell of the three-dimensional metamaterial structure are depicted in Figs. 1(a) and 1(b). The arrayed 3D structures consist of standup DSRRs with gaps on the top and bottom, forming two capacitors connected in series in an equivalent  $LC$  circuit. Silicon, the photoactive material in these structures, is incorporated into the bottom gap of the DSRR. This structure is resonant when the THz wave is normally incident on the sample with the  $E$  field parallel to the sides with the gaps ( $H$  then passes through the DSRRs). In the absence of photoexcitation, the THz pulses couple to the resonators magnetically, inducing a circulating current in the ring. In particular, the symmetry is such that the electric field does not couple to the DSRRs (except for substrate-induced symmetry breaking, as discussed below). Upon optical excitation of the silicon, free carriers are generated, resulting in an increased conductivity. The conducting silicon gradually shorts the capacitance of the lower gap, which, in turn, increases the total capacitance, resulting in a redshift of the  $LC$  resonance. With sufficient pump power, the bottom capacitor can be effectively shorted, leading to the maximum possible frequency shift. Importantly, with the lower gap shorted, the symmetry of the DSRR is reduced such that the incident  $E$  field now drives circulating

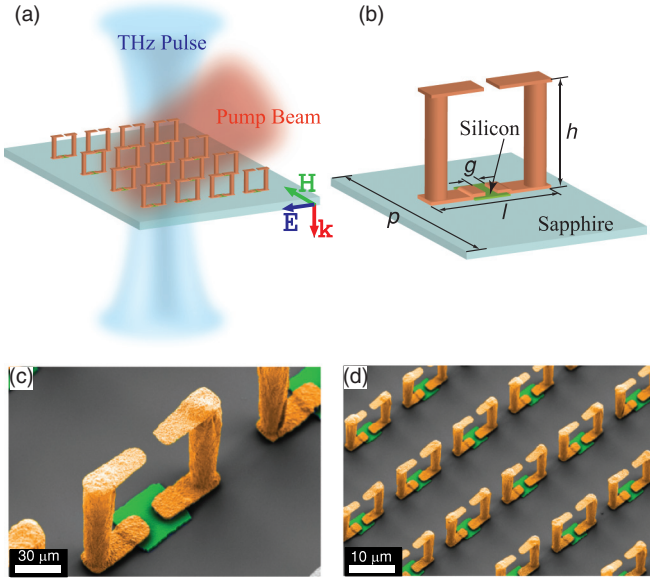


FIG. 1. (Color online) Schematic of the design of the broadband tunable three-dimensional metamaterials at THz frequencies and scanning electron micrographs (SEMs) of fabricated 3D structures. (a) Schematic principle of the photoexcitation tuning of the 3D metamaterials. (b) Typical dimensions for unit cell of DSRRs:  $p = 50 \mu\text{m}$ ,  $h = 33 \mu\text{m}$ ,  $g = 5 \mu\text{m}$ , and  $l = 36 \mu\text{m}$ . (c) The oblique closeup view of a fabricated unit cell of DSRRs in copper standing on a sapphire substrate. (d) Oblique view of arrayed hybrid 3D metamaterials.

currents in the resonator (in addition to the  $H$  field). That is, photoexcitation induces intrinsic bianisotropy.

Multilayer electroplating (MLEP) in combination with conventional optical lithography is utilized to fabricate the 3D metamaterials on a commercial silicon-on-sapphire wafer consisting of 600-nm (100) intrinsic silicon on top of a 530- $\mu\text{m}$ -thick  $R$ -plane sapphire substrate. The fabrication of 3D DSRRs begins with patterning of the distributed silicon pads using reactive ion etching. Then two 1.5- $\mu\text{m}$ -thick bottom copper laminas were electroplated on a copper seed layer, forming a 5- $\mu\text{m}$ -wide gap between the two laminas. The laminas were patterned along the primary flat orientation of the wafer, which is  $45^\circ$  counterclockwise from the projection of the  $C$  axis on the  $R$  plane. Thus, the refractive indices on two axes are the same. The patterned silicon pads sit beneath the gap. After removing the photoresist, a 30- $\mu\text{m}$ -thick layer of photoresist (PR) AZ9260 was patterned with two 6- $\mu\text{m}$ -diam holes at both ends of the laminas to enable electroplating. Without removing the thick PR, another 100-nm seed layer of copper was evaporated on the top without any wrinkling. After one thin layer PR was patterned, 1.5- $\mu\text{m}$  laminas that are the same as the bottom ones were fabricated to connect the standing pillars. As a final releasing step, the PR was removed with acetone, and the copper seed layers are etched away in dilute acid. Using MLEP, the height of the DSRRs can be determined by the thickness of PR, which is controlled by the spinning rate. Figures 1(c) and 1(d) show the SEM images of the fabricated 3D metamaterials sample.

Terahertz time-domain spectroscopy (THz-TDS) was used to characterize the frequency-dependent electromagnetic

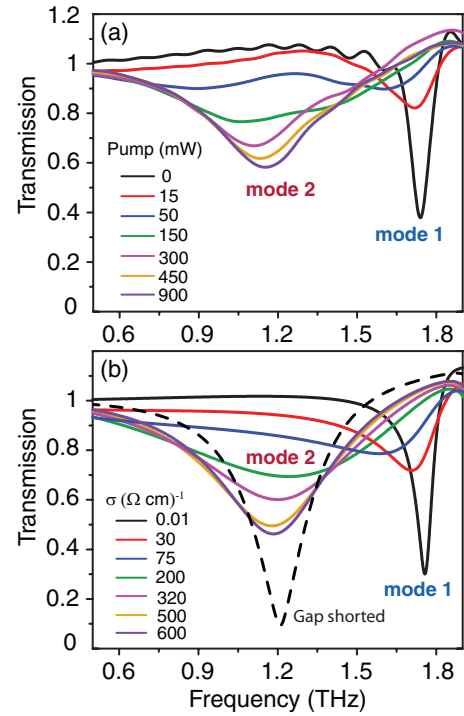


FIG. 2. (Color online) (a) Experimentally measured frequency-dependent THz electric field transmission. The response is shown for increasing laser excitation power. (b) Full wave electromagnetic simulation results of the transmission vs frequency for various values of the silicon conductivity. The dashed black curve is a simulation with the bottom gap shorted with copper.

response of the photoactive 3D DSRRs. A bare sapphire substrate with the same thickness as the sample substrate was used as the reference. The sample and reference were mounted on the holder with the same orientation as to minimize sapphire birefringence. The THz pulses were at normal incidence to the substrate with  $E$  field parallel to the side of the DSRRs with the gaps [see Fig. 1(a)]. Photoexcitation of free carriers in the silicon was achieved with 1.55-eV, 35-fs ultrafast pulses at a repetition rate of 1 kHz. The optical pump pulse was set to arrive 10 ps before the THz probe beam to achieve a near steady-state accumulation of carriers due to their long lifetime in silicon. To ensure homogeneous excitation, the pump laser beam was 8 mm in diameter while the incident THz had a 3 mm diameter.

The experimental results are shown in Fig. 2(a). The black curve shows the response without photoexcitation. There is a very sharp resonance at 1.74 THz ( $\omega_1$ ) with a minimum transmission of 35%. As the pump power is increased to 450 mW, a pronounced resonance shift to 1.16 THz ( $\omega_2$ ) is observed with a transmission of 60%. Further, a tunability of 30% of the resonance frequency is achieved. At higher powers, a large density of carriers result in significant carrier-carrier scattering, saturating the conductivity. For example, increasing the excitation power from 450 to 900 mW (corresponding to a fluence of  $1.8 \text{ mJ cm}^{-2}$ ) leads to only a minor change in the DSRR electromagnetic response.

Full wave electromagnetic simulations were performed using CST microwave studio. To compare with experiments, the conductivity of the silicon comprising the substrate capacitor was varied in the simulations. The photoconductive silicon is

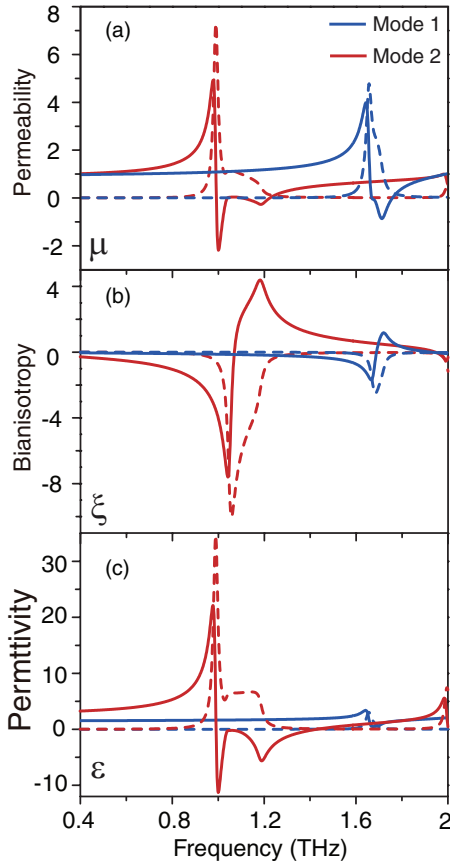


FIG. 3. (Color online) Retrieved constitutive parameters of (a) permeability  $\mu$ , (b) bianisotropy parameter  $\xi$ , and (c) permittivity  $\epsilon$ , from full wave electromagnetic simulations of the 3D DSRRs. The blue lines represent the parameters at mode 1 (i.e., no photoexcitation). The red lines represent the parameters at mode 2 assuming the bottom gap is shorted by replacing the silicon by copper. Solid lines are real parts and dashed lines are imaginary parts.

simulated as dielectric with  $\epsilon_{\text{Si}} = 11.7$  and a pump-power-dependent conductivity of  $\sigma_{\text{Si}}$ . The substrate sapphire was modeled as lossless dielectric with  $\epsilon_{\text{sapphire}} = 10.5$ . The copper was modeled as a lossy metal with a frequency-independent conductivity of  $5 \times 10^4 (\Omega \text{ cm})^{-1}$ , based on four-point probe measurements of an electroplated continuous Cu film. The solid black curve in Fig. 3(b) corresponds to 0 mW pump power with  $\sigma_{\text{Si}}$  taken as  $0.01 (\Omega \text{ cm})^{-1}$ , while for illumination of 450 mW (fluence of  $0.9 \text{ mJ cm}^{-2}$ ), the conductivity is estimated as  $500 (\Omega \text{ cm})^{-1}$ , which is close to the reported value in Ref. 13, given the variability of the properties of the silicon grown on sapphire, the subsequent fabrication process, etc. As shown in Fig. 2(b), the simulated and measured frequencies match each other quite well, highlighting that the observed changes in the electromagnetic response arise from changes in the Si conductivity.

For our experiments, the electric field is normal to the gaps and the magnetic field is normal to the plane of rings. As such, the electric and magnetic components of the THz waves can, in principle, couple to the DSRRs leading to a bianisotropic response. However, without photoexcitation (mode 1), inversion symmetry of the ring (in the propagation direction) eliminates the possibility of electric field excitation resulting

in a pure magnetic response (the substrate breaks this inversion symmetry and leads to measurable, but small, bianisotropy—see below and Fig. 3). Following photoexcitation, the shorted bottom gap breaks the inversion symmetry, leading to strong electric field excitation and a bianisotropic response.

To further analyze the electromagnetic response of modes 1 and 2, the real and imaginary parts of  $\epsilon$ ,  $\mu$  and  $\xi$ , were extracted from simulations using a bianisotropic retrieval method.<sup>23–26</sup> As shown in Fig. 3(b), at mode 1, the permeability exhibits a strong Lorentzian-like response at the resonance yielding negative values in a certain span. However, as mentioned above, the substrate breaks the symmetry and induces a small bianisotropy as the blue curves in Fig. 3(b) indicate.<sup>27</sup> Hence, there is also a small resonant response for the permittivity at mode 1. When the bottom gap is totally shorted (mode 2), the 3D DSRR array becomes fully bianisotropic with both the electric and magnetic field resonantly driving the array. The bianisotropic response is now dominated by the DSRRs. Although the permittivity and permeability present negative values, the refractive index is still positive due to the strong bianisotropy. Nonetheless, our metamaterials can be used to effectively switch the DSRRs with the electromagnetic response changing from near purely magnetic to bianisotropic.

From the measurements and simulations, the mode 2 resonance is broadened in comparison to mode 1. While there are losses from excited carriers in the silicon, this is not the dominant source of the broadening. This is evident in Fig. 2(b), showing that the simulation with the lower gap shorted with copper (black dashed line) is still considerably broader than the mode 1 resonance. In fact, the broadening of mode 2 results from increased radiation damping.<sup>28</sup> Additional insight into the electromagnetic response and the nature of this damping is obtained through consideration of the local electric field and current density (from simulations) at the unit cell level. Figure 4 shows the simulated surface currents and the electric field normal to the gap at both modes respectively. We note that the peak of the surface currents is  $90^\circ$  out of phase with the peak of electric field. For mode 1, the circulating current causes

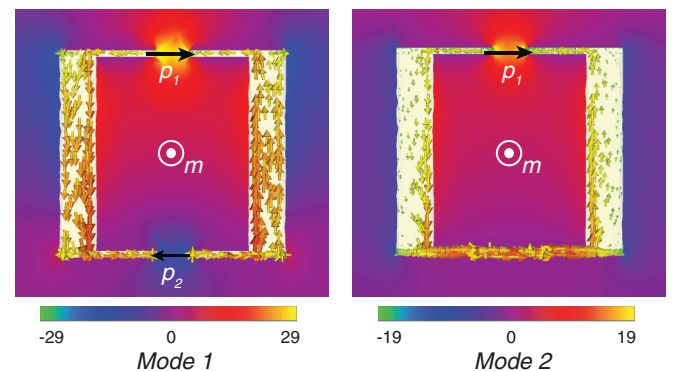


FIG. 4. (Color online) Simulation of the distribution of surface current density and electric field in the direction of normal to the gaps at two modes. Left: Surface current and electric field at mode 1 with a magnetic dipole of  $m$  and electric dipoles of  $p_1$  and  $p_2$ . Right: Surface current and electric field at mode 2 with a magnetic dipole of  $m$  and an electric dipoles of  $p_1$ . The electric field and the surface current are  $90^\circ$  out of phase. The color bars represent the simulated excited field strength relative to the incident field.



charge accumulation at both gaps forming two electric dipoles  $p_1$  and  $p_2$  in opposite directions, which can be equivalently taken as a quadrupole as shown on the left side of Fig. 4. As such, radiation damping of the DSRRs for mode 1 is at the level of magnetic dipole and electric quadrupole terms in a multipole expansion of the electrodynamic response. In contrast, for mode 2 (in which the bottom gap is effectively shorted), there is only the top gap, which is equivalent to an electric dipole  $p_1$  (right side, Fig. 4). Consequently, radiation damping for mode 2 is dominated by the electric dipole contribution, leading to a decrease in the quality factor of the resonance. It is important to note that this is intrinsic to the SRRs and is not a spurious contribution that can, for our geometry, be minimized.

In summary, we have successfully fabricated and characterized broadband tunable 3D hybrid metamaterials at THz frequencies. With photoexcitation of silicon, we observed over

30% redshift of the resonant frequency, in agreement with numerical simulations. Parameters retrieval indicates that our 3D structure can be effectively switched from a mode with a strong magnetic response to a fully bianisotropic response. Our tunable 3D metamaterials can be potentially used as a tunable notch filter, high speed modulator. With proper design on the phase shift between two modes, artificial polarizers can also be realized.

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<sup>1</sup>D. Schurig, J. J. Mock, B. J. Justice, S. A. Cummer, J. B. Pendry, A. F. Starr, and D. R. Smith, *Science* **314**, 977 (2006).

<sup>2</sup>J. Valentine, J. Li, T. Zentgraf, G. Bartal, and X. Zhang, *Nat. Mater.* **8**, 568 (2009).

<sup>3</sup>T. Ergin, N. Stenger, P. Brenner, J. B. Pendry, and M. Wegener, *Science* **328**, 337 (2010).

<sup>4</sup>R. A. Shelby, D. R. Smith, and S. Schultz, *Science* **292**, 77 (2001).

<sup>5</sup>J. Valentine, S. Zhang, T. Zentgraf, E. Ulin-Avila, D. A. Genov, G. Bartal, and X. Zhang, *Nature (London)* **455**, 376 (2008).

<sup>6</sup>R. Marquès, F. Medina, and R. Rafii-El-Idrissi, *Phys. Rev. B* **65**, 144440 (2002).

<sup>7</sup>N. I. Landy, S. Sajuyigbe, J. J. Mock, D. R. Smith, and W. J. Padilla, *Phys. Rev. Lett.* **100**, 207402 (2008).

<sup>8</sup>H. Tao, N. I. Landy, C. M. Bingham, X. Zhang, R. D. Averitt, and W. J. Padilla, *Opt. Express* **16**, 7181 (2008).

<sup>9</sup>X. Liu, T. Starr, A. F. Starr, and W. J. Padilla, *Phys. Rev. Lett.* **104**, 207403 (2010).

<sup>10</sup>H.-T. Chen, W. J. Padilla, J. M. O. Zide, A. C. Gossard, A. J. Taylor, and R. D. Averitt, *Nature (London)* **444**, 597 (2006).

<sup>11</sup>H.-T. Chen, W. J. Padilla, M. J. Cich, A. K. Azad, R. D. Averitt, and A. J. Taylor, *Nat. Photonics* **3**, 148 (2009).

<sup>12</sup>W. J. Padilla, A. J. Taylor, C. Highstrete, M. Lee, and R. D. Averitt, *Phys. Rev. Lett.* **96**, 107401 (2006).

<sup>13</sup>H.-T. Chen, J. F. O'Hara, A. K. Azad, A. J. Taylor, R. D. Averitt, D. B. Shrekenhamer, and W. J. Padilla, *Nat. Photonics* **2**, 295 (2008).

<sup>14</sup>N. H. Shen, M. Massouti, M. Gokkavas, J.-M. Manceau, E. Ozbay, M. Kafesaki, T. Koschny, S. Tzortzakis, and C. M. Soukoulis, *Phys. Rev. Lett.* **106**, 037403 (2011).

<sup>15</sup>T. Driscoll, S. Palit, M. M. Qazilbash, M. Brehm, F. Keilmann, B.-G. Chae, S.-J. Yun, H.-T. Kim, S. Y. Cho, N. M. Jokerst, D. R. Smith, and D. N. Basov, *Appl. Phys. Lett.* **93**, 024101 (2008).

<sup>16</sup>H.-T. Chen, H. Yang, R. Singh, J. F. O'Hara, A. K. Azad, S. A. Trugman, Q. X. Jia, and A. J. Taylor, *Phys. Rev. Lett.* **105**, 247402 (2010).

<sup>17</sup>J. Zhu, J. Han, Z. Tian, J. Gu, Z. Chen, and W. Zhang, *Opt. Commun.* **284**, 3129 (2011).

<sup>18</sup>H. Tao, A. C. Strikwerda, K. Fan, W. J. Padilla, X. Zhang, and R. D. Averitt, *Phys. Rev. Lett.* **103**, 147401 (2009).

<sup>19</sup>W. M. Zhu, A. Q. Liu, X. M. Zhang, D. P. Tsai, T. Bourouina, J. H. Teng, X. H. Zhang, H. C. Guo, H. Tanoto, T. Mei, G. Q. Lo, and D. L. Kwong, *Adv. Mater.* **23**, 1792 (2011).

<sup>20</sup>S. Zhang, Y.-S. Park, J. Li, X. Lu, W. Zhang, and X. Zhang, *Phys. Rev. Lett.* **102**, 023901 (2009).

<sup>21</sup>K. Fan, A. C. Strikwerda, H. Tao, X. Zhang, and R. D. Averitt, *Opt. Express* **19**, 12619 (2011).

<sup>22</sup>S. Zhang, J. Zhou, Y.-S. Park, J. Rho, R. Singh, S. Nam, A. K. Azad, H.-T. Chen, X. Yin, A. J. Taylor, and X. Zhang, *Nat. Commun.* **3**, 942 (2012).

<sup>23</sup>R. Liu, T. J. Cui, D. Huang, B. Zhao, and D. R. Smith, *Phys. Rev. E* **76**, 026606 (2007).

<sup>24</sup>C. E. Kriegler, M. S. Rill, S. Linden, and M. Wegener, *IEEE J. Sel. Top. Quantum Electron.* **16**, 367 (2010).

<sup>25</sup>R. Zhao, T. Koschny, and C. M. Soukoulis, *Opt. Express* **18**, 14553 (2010).

<sup>26</sup>D. R. Smith, *Phys. Rev. E* **81**, 036605 (2010).

<sup>27</sup>D. A. Powell and Y. S. Kivshar, *Appl. Phys. Lett.* **97**, 091106 (2010).

<sup>28</sup>T. Q. Li, H. Liu, T. Li, S. M. Wang, J. X. Cao, Z. H. Zhu, Z. G. Dong, S. N. Zhu, and X. Zhang, *Phys. Rev. B* **80**, 115113 (2009).